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NEWS 1		Web Page for STN Seminar Schedule - N. America
NEWS 2	NOV 21	CAS patent coverage to include exemplified prophetic substances identified in English-, French-, German-, and Japanese-language basic patents from 2004-present
NEWS 3	NOV 26	MARPAT enhanced with FSORT command
NEWS 4	NOV 26	CHEMSAFE now available on STN Easy
NEWS 5	NOV 26	Two new SET commands increase convenience of STN searching
NEWS 6	DEC 01	ChemPort single article sales feature unavailable
NEWS 7	DEC 12	GBFULL now offers single source for full-text coverage of complete UK patent families
NEWS 8	DEC 17	Fifty-one pharmaceutical ingredients added to PS
NEWS 9	JAN 06	The retention policy for unread STNmail messages will change in 2009 for STN-Columbus and STN-Tokyo
NEWS 10	JAN 07	WPIDS, WPINDEX, and WPIX enhanced Japanese Patent Classification Data
NEWS 11	FEB 02	Simultaneous left and right truncation (SLART) added for CERAB, COMPUAB, ELCOM, and SOLIDSTATE
NEWS 12	FEB 02	GENBANK enhanced with SET PLURALS and SET SPELLING
NEWS 13	FEB 06	Patent sequence location (PSL) data added to USGENE
NEWS 14	FEB 10	COMPENDEX reloaded and enhanced
NEWS 15	FEB 11	WTEXTILES reloaded and enhanced
NEWS 16	FEB 19	New patent-examiner citations in 300,000 CA/CAplus patent records provide insights into related prior art
NEWS 17	FEB 19	Increase the precision of your patent queries -- use terms from the IPC Thesaurus, Version 2009.01
NEWS 18	FEB 23	Several formats for image display and print options discontinued in USPATFULL and USPAT2
NEWS 19	FEB 23	MEDLINE now offers more precise author group fields and 2009 MeSH terms
NEWS 20	FEB 23	TOXCENTER updates mirror those of MEDLINE - more precise author group fields and 2009 MeSH terms
NEWS 21	FEB 23	Three million new patent records blast AEROSPACE into STN patent clusters
NEWS 22	FEB 25	USGENE enhanced with patent family and legal status display data from INPADOCDB
NEWS EXPRESS		JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3, AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.
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* * * * * * * * * * * * * * * STN  Columbus  * * * * * * * * * * * * * * *

FILE 'HOME' ENTERED AT 14:30:02 ON 01 MAR 2009

=> file reg
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
FULL ESTIMATED COST                                     0.22       0.22

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DICTIONARY FILE UPDATES:  27 FEB 2009  HIGHEST RN 1113101-98-6

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=>
Uploading C:\Program Files\Stnexp\Queries\10579106restriction.str
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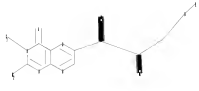

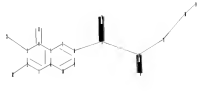

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ring nodes :
1 2 3 4 5 6 7 8 9 10
chain bonds :
2-12 3-21 4-11 8-13 13-14 13-15 15-16 15-17 17-18 18-19
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-10 7-8 8-9 9-10
exact/norm bonds :
1-2 1-6 2-3 2-12 3-4 3-21 4-5 4-11 13-14 15-16 17-18
exact bonds :
8-13 13-15 15-17 18-19
normalized bonds :
5-6 5-7 6-10 7-8 8-9 9-10
isolated ring systems :
containing 1 :

G1:C,H

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom
11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:CLASS 18:CLASS
19:CLASS 21:CLASS

L1 STRUCTURE UPLOADED

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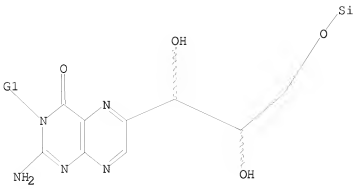

G1 C,H

Structure attributes must be viewed using STN Express query preparation.

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SAMPLE SCREEN SEARCH COMPLETED -        2 TO ITERATE

100.0% PROCESSED         2 ITERATIONS                                0 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:  ONLINE   **COMPLETE**
                        BATCH    **COMPLETE**
PROJECTED ITERATIONS:             2 TO      124
PROJECTED ANSWERS:                0 TO        0

L2            0 SEA SSS SAM L1

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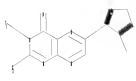

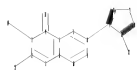

chain nodes :
11 12 14 21
ring nodes :
1 2 3 4 5 6 7 8 9 10 16 17 18 19 20
chain bonds :
2-12 3-14 4-11 8-17 16-21
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-10 7-8 8-9 9-10 16-17 16-20 17-18 18-19
19-20
exact/norm bonds :
1-2 1-6 2-3 2-12 3-4 3-14 4-5 4-11 16-17 16-20 17-18 18-19 19-20
exact bonds :
8-17 16-21
normalized bonds :
5-6 5-7 6-10 7-8 8-9 9-10
isolated ring systems :
containing 1 :

G1:C,H

G2:S,N

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom
11:CLASS 12:CLASS 14:CLASS 16:Atom 17:Atom 18:Atom 19:Atom 20:Atom 21:CLASS

L3 STRUCTURE UPLOADED

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L3 HAS NO ANSWERS
L3                STR
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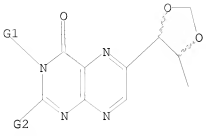

G1 C,H

G2 S,N

Structure attributes must be viewed using STN Express query preparation.

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=> s l3 full
FULL SEARCH INITIATED 14:34:07 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -       19 TO ITERATE

100.0% PROCESSED       19 ITERATIONS                                 6 ANSWERS
SEARCH TIME: 00.00.01

L4             6 SEA SSS FUL L3

=> file caplus
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
FULL ESTIMATED COST                                   188.76     188.98

FILE 'CAPLUS' ENTERED AT 14:34:11 ON 01 MAR 2009
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FILE COVERS 1907 - 1 Mar 2009  VOL 150 ISS 10
FILE LAST UPDATED: 27 Feb 2009  (20090227/ED)

Caplus now includes complete International Patent Classification (IPC)
reclassification data for the third quarter of 2008.
```

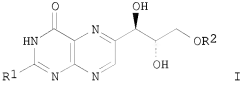
=> s l4 full
L5 5 L4

=> d ibib abs hitstr tot

L5 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2005:472157 CAPLUS
DOCUMENT NUMBER: 143:7534
TITLE: Preparation of tetrahydrobiopterin and analogs of tetrahydrobiopterin
INVENTOR(S): Moser, Rudolf; Groehn, Viola; Schumacher, Andreas; Martin, Pierre
PATENT ASSIGNEE(S): Biomarin Pharmaceutical Inc., USA; Merck Eprova A.-G.
SOURCE: PCT Int. Appl., 55 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005049614	A2	20050602	WO 2004-US38313	20041117
WO 2005049614	A3	20070308		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2004290692	A1	20050602	AU 2004-290692	20041117
CA 2545484	A1	20050602	CA 2004-2545484	20041117
EP 1776364	A2	20070425	EP 2004-819154	20041117
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, HR, LT, LV, MK, YU				
JP 2007534637	T	20071129	JP 2006-539994	20041117
US 20070244322	A1	20071018	US 2007-579106	20070216
PRIORITY APPLN. INFO.:			US 2003-520367P	P 20031117
			US 2003-520368P	P 20031117
			WO 2004-US38313	W 20041117

OTHER SOURCE(S): CASREACT 143:7534; MARPAT 143:7534
GI



I

AB A process for the preparation of tetrahydrobiopterin and its analogs, e.g. I [R1 = alkylamino, arylamino, alkylthio, alkylaminomethyleneimino, R2 = H; R1 = alkylamino, alkylthio, Me2NCH:N, R2 = Me2CHEt2Si, (Me3CO)Ph2Si,

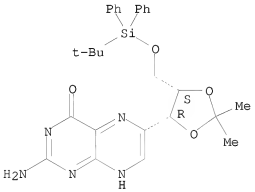
MePh2Si, Me3CMe2Si, Me3C(MeO)PhSi, (Me3C)2MeSi, etc.], from neopterin and/or 6-substituted pterins with an improved yield and a high stereoselectivity is disclosed. Also disclosed herein are novel individual intermediates prepared in the preparation of tetrahydrobiopterin, such as selectively protected neopterin useful for the preparation of tetrahydrobiopterin. As an example, L-neopterin was reacted with DMF-acetal to give the 2-(dimethylamino)methylene derivative I (R1 = Me2NCH:N, R2 = H) (II). II was then silylated to I (R2 = Me3CPh2Si) which could be deprotected to I (R1 = NH2).

IT 852547-48-9P
RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)
(preparation of tetrahydrobiopterin and analogs)

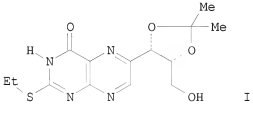
RN 852547-48-9 CAPLUS

CN 4(3H)-Pteridinone, 2-amino-6-[(4R,5S)-5-[[[(1,1-dimethylethyl)diphenylsilyl]oxy]methyl]-2,2-dimethyl-1,3-dioxolan-4-yl]- (CA INDEX NAME)

Absolute stereochemistry.



L5 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2000:499841 CAPLUS
DOCUMENT NUMBER: 133:207729
TITLE: Synthesis of 2-ethylthio-6-(3-hydroxy-1,2-O-isopropylidenepropyl)pteridin-4(3H)-one
AUTHOR(S): Kang, Yonghan; Kim, Seungjin; Myoung, Youngchan; Baek, Daejin
CORPORATE SOURCE: Department of Chemistry, Hanyang University, Ansan, 425-791, S. Korea
SOURCE: Heterocycles (2000), 53(7), 1551-1557
CODEN: HTCYAM; ISSN: 0385-5414
PUBLISHER: Japan Institute of Heterocyclic Chemistry
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 133:207729
GI



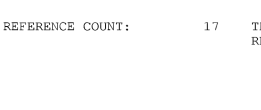
I

AB A strategy has been described for the synthesis of 2-ethylthio-6-(3-hydroxy-1,2-O-isopropylidenepropyl)pteridin-4(3H)-one (I), which can be used as a useful intermediate for the conversion of neopterin to biopterin.

IT 290370-92-2P
RL: SPN (Synthetic preparation); PREP (Preparation)
(synthesis of 2-ethylthio-6-(3-hydroxy-1,2-O-isopropylidenepropyl)pteridin-4(3H)-one, a useful intermediate in the synthesis of biopterin)

RN 290370-92-2 CAPLUS
CN 4(3H)-Pteridinone, 2-(ethylthio)-6-[(4S,5R)-5-(hydroxymethyl)-2,2-dimethyl-1,3-dioxolan-4-yl]- (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 17 THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1995:536893 CAPLUS
DOCUMENT NUMBER: 122:314347
ORIGINAL REFERENCE NO.: 122:57165a,57168a
TITLE: Pteridines CV. Selective N(3)- and O4-alkylation of L-biopterin: A convenient synthesis of 3- and O4-methyl-L-biopterin and the versatile N2-(N,N-dimethylaminomethylene)-N(3)-p-nitrophenethyl-protected L-biopterin
AUTHOR(S): Hanaya, Tadashi; Torigoe, Kiyoshi; Soranaka, Kazuyuki; Yamamoto, Horoshi; Qizheng, Yao; Pfleiderer, Wolfgang
CORPORATE SOURCE: Faculty Science, Okayama University, Okayama, 700, Japan
SOURCE: Pteridines (1995), 6(1), 1-7
CODEN: PTRDEO; ISSN: 0933-4807
PUBLISHER: International Society of Pteridinology
DOCUMENT TYPE: Journal
LANGUAGE: English
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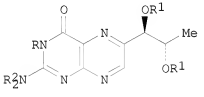


I

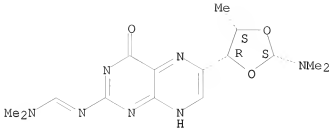
AB Treatment of L-biopterin with N,N-dimethylformamide dimethyl- (or diethyl)acetal and then with acetic anhydride in pyridine gave 1',2'-di-O-acetyl-N2-(N,N-dimethylaminoethylene)-L-biopterin, which was converted by the Mitsunobu reaction into 3-Me and 3-p-nitrophenetyl derivs. The protective groups on the side chain diols and N2 of these compds. were selectively cleaved to furnish biopterins I (R = Me, R1 = H, R22 = CHNMe2; R = 4-O2NC6H4CH2CH2, R1 = H, R22 = CHNMe2; R = Me, 4-O2NC6H4CH2CH2, R1 = R2 = H), among which I (R = Me, R1 = R2 = H) is naturally occurring 3-methyl-L-biopterin and I (R = 4-O2NC6H4CH2CH2, R1 = H, R22 = CHNMe2) is N2,N(3)-protected biopterin, a versatile intermediate for various reactions on the side-chain diol. In contrast, the same Mitsunobu reactions of tri-N2:1',2'-O-acetyl-L-biopterin afforded O4-Me and O4-NPE derivs., both of which yielded O4-methyl-L-biopterin and subsequently led to 4-amino-L-biopterin.

IT 163132-77-2P 163252-46-8P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of alkyl biopterins)

RN 163132-77-2 CAPLUS

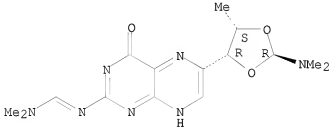
CN Methanimidamide, N'-[6-[2-(dimethylamino)-5-methyl-1,3-dioxolan-4-yl]-3,4-dihydro-4-oxo-2-pteridinyl]-N,N-dimethyl-, [2S-(2α,4α,5α)]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.



RN 163252-46-8 CAPLUS
CN Methanimidamide, N'-[6-[2-(dimethylamino)-5-methyl-1,3-dioxolan-4-yl]-3,4-dihydro-4-oxo-2-pteridinyl]-N,N-dimethyl-, [2R-(2α,4β,5β)]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.



L5 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1990:55443 CAPLUS
DOCUMENT NUMBER: 112:55443
ORIGINAL REFERENCE NO.: 112:9511a,9514a
TITLE: Synthesis of (-)-biopterin using (S)-ethyl lactate as a starting material
AUTHOR(S): Kikuchi, Haruhiko; Mori, Kenji
CORPORATE SOURCE: Dep. Agric. Chem., Univ. Tokyo, Tokyo, 113, Japan
SOURCE: Agricultural and Biological Chemistry (1989), 53(8), 2095-100
CODEN: ABCHA6; ISSN: 0002-1369
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 112:55443
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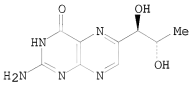
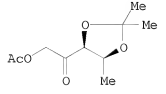


I



III

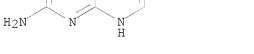
AB (-)-Biopterin (I) was synthesized from (1S,2S)-1-(1,3-dithian-2-yl)propane-1,2-diol (II), which was derived from com. available Et (S)-lactate. II was converted to ketone III through a six-step sequence. III was submitted to condensation with 3,5,6-triaminopyrimidinol, and followed by oxidation to afford isopropylidenebiopterin which was deprotected to give I.

IT 124600-40-4P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and deblocking of)

RN 124600-40-4 CAPLUS

CN 4(1H)-Pteridinone, 2-amino-6-(2,2,5-trimethyl-1,3-dioxolan-4-yl)-, (4R-cis)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L5 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 1972:126936 CAPLUS
DOCUMENT NUMBER: 76:126936
ORIGINAL REFERENCE NO.: 76:20553a,20556a
TITLE: Pterin chemistry. 41. New synthesis of DL-biopterin
AUTHOR(S): Viscontini, M.; Frei, W. F.
CORPORATE SOURCE: Org.-Chem. Inst., Univ. Zurich, Zurich, Switz.
SOURCE: Helvetica Chimica Acta (1972), 55(2), 574-9
CODEN: HCACAV; ISSN: 0018-019X
DOCUMENT TYPE: Journal
LANGUAGE: German

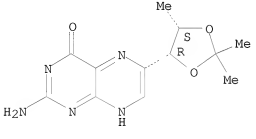
AB D,L-Biopterin was prepared in .apprx.40% yield by condensing D,L-2,3-O-isopropylidene-4-methylerythrulose (I) with 2,4,5-triamino-6-hydroxypyrimidine, and oxidation with air to give D,L-1',2'-O-isopropylidenebiopterin. The isopropylidene protective group was easily hydrolyzed off to give isomer-free D,L-biopterin. I was prepared by oxidizing trans-crotonic acid to D,L-erythro-2,3-dihydroxybutyric acid, and introducing the protective group before the extra C.

IT 36183-32-1P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

RN 36183-32-1 CAPLUS

CN 4(3H)-Pteridinone, 2-amino-6-[(4R,5S)-2,2,5-trimethyl-1,3-dioxolan-4-yl]-, rel- (CA INDEX NAME)

Relative stereochemistry.



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=> FIL STNGUIDE
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
FULL ESTIMATED COST                                    29.20     218.18

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)          SINCE FILE      TOTAL
                                                       ENTRY    SESSION
CA SUBSCRIBER PRICE                                    -4.10      -4.10

FILE 'STNGUIDE' ENTERED AT 14:35:22 ON 01 MAR 2009
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FILE CONTAINS CURRENT INFORMATION.
LAST RELOADED: Feb 27, 2009 (20090227/UP).

=> d his

     (FILE 'HOME' ENTERED AT 14:30:02 ON 01 MAR 2009)

     FILE 'REGISTRY' ENTERED AT 14:30:16 ON 01 MAR 2009
L1               STRUCTURE UPLOADED
L2            0 S L1
L3               STRUCTURE UPLOADED
L4            6 S L3 FULL

     FILE 'CAPLUS' ENTERED AT 14:34:11 ON 01 MAR 2009
L5            5 S L4 FULL

     FILE 'STNGUIDE' ENTERED AT 14:35:22 ON 01 MAR 2009

=> log y
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
FULL ESTIMATED COST                                     0.63     218.81

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)          SINCE FILE      TOTAL
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CA SUBSCRIBER PRICE                                     0.00      -4.10

STN INTERNATIONAL LOGOFF AT 14:40:57 ON 01 MAR 2009
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